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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/075,970	02/14/2002	Michael Helmus	01-202 (4010/15)	9278
27774 7590 08/05/2011 MAYER & WILLIAMS PC 251 NORTH AVENUE WEST Suite 201 WESTFIELD, NJ 07090			EXAMINER TYSON, MELANIE RUANO	
			ART UNIT 3773	PAPER NUMBER
			MAIL DATE 08/05/2011	DELIVERY MODE PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary

Application No.

10/075,970

Applicant(s)

HELMUS, MICHAEL

Examiner

MELANIE TYSON

Art Unit

3773

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --
Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 21 June 2011.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1,3,5-7,9-15,17,19-21 and 48-50 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 1,3,5-7,9-15,17,19-21 and 48-50 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
- ☐ Certified copies of the priority documents have been received.
 - ☐ Certified copies of the priority documents have been received in Application No. _____.
 - ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- 1) ☒ Notice of References Cited (PTO-892)
- 2) ☐ Notice of Draftperson's Patent Drawing Review (PTO-946)
- 3) ☐ Information Disclosure Statement(s) (PTO/SB08)
Paper No(s)/Mail Date _____
- 4) ☐ Interview Summary (PTO-413)
Paper No(s)/Mail Date _____
- 5) ☐ Notice of Informal Patent Application
- 6) ☐ Other: _____

DETAILED ACTION

This action is in response to the applicant's amendment received 21 June 2011. The amendments made to the claims do not place the application in condition for allowance for the reasons set forth below. Claims 2, 4, 8, 16, 18, and 22-47 are cancelled.

Response to Arguments

Applicant's arguments with respect to the claims have been considered but are moot in view of the new ground(s) of rejection.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

The factual inquiries set forth in *Graham v. John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

1. Determining the scope and contents of the prior art.
2. Ascertaining the differences between the prior art and the claims at issue.
3. Resolving the level of ordinary skill in the pertinent art.
4. Considering objective evidence present in the application indicating obviousness or nonobviousness.

Claims 1, 3, 5-7, 9-15, 17, 19-21, and 48-50 are rejected under 35 U.S.C. 103(a) as being unpatentable over Hossainy et al. (U.S. Patent No. 6,153,252) in

view of Bolz (U.S. Patent No. 6,287,332 B1) and Igaki et al. (U.S. Patent No. 6,045,568).

Hossainy discloses an implantable medical device (stent; see entire document) comprising a biodegradable inner core (for example, see column 3, lines 10-21), thus becoming decreasingly rigid upon contact with bodily fluid, a biodegradable covering material completely covering the inner core material as a coating thereon (for example, see column 3, lines 55-58 and column 4, lines 1-14) and does not contain a therapeutic agent therein (an alternate embodiment may contain agents if desired), and one or more coating layers that may be provided on the inner core material and/or the covering material that do not contain therapeutic agents (for example, see column 7, lines 11-15), wherein the entire medical device is substantially biodegradable by the body (i.e., both the cover and inner core may be biodegradable). The covering material may be formed of a hydrophobic surface erodable polymer (for example, polyamide, polyorthoester, or polyanhydride; for example, see column see 5, lines 6-12), thus is capable of controlling the rate at which the inner core material becomes flexible upon contact with bodily fluids. Hossainy discloses the inner core may be metallic, an absorbable plastic, or any other suitable material which can provide the necessary mechanical requirements of a stent, but fails to disclose the biodegradable inner core material is specifically selected from biodegradable metallic or ceramic materials.

Bolz discloses an implantable medical device, such as a bioresorbable stent (see entire document). Bolz teaches constructing the bioresorbable stent of degradable metallic materials. Bolz further teaches that stents of degradable metallic material

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combine the advantageous mechanical properties of metal stents (such as elasticity, deformability, and stability by way of improving ductility, tensile strength, etc.; for example, see column 3, lines 11-35) with the bioresorbability of polymer-based stents (for example, see column 2, lines 6-16). Therefore, it would have been obvious to one of ordinary skill in the art at the time the invention was made to form Hossainy's inner core from a biodegradable metallic material as taught by Bolz. Doing so would provide the mechanical advantages described above. Hossainy as modified by Bolz fails to disclose or suggest the medical device maintains adequate rigidity for a period from about three to six months following implantation.

Igaki discloses an implantable medical device, such as a bioresorbable stent (see entire document). Igaki teaches the bioresorbable stent substantially maintains its shape until after about three to six months of implantation and is absorbed in the body in about six to twelve months (for example, see experiment 1). Igaki suggests this range is adequate to prevent reconstriction of the blood vessel without inflammation or excess hypertrophy of the vessel (for example, see column 5, lines 61-67). Hossainy recognizes absorption rates of the polymeric covering materials may be adjusted as needed (for example, see column 7, lines 18-55). Bolz also recognizes stents for some applications are only needed for a few months, afterwards the part of the vessel affected by, for example, stenosis would remain open without any support (for example, see column 1, lines 38-42). Therefore, would have been obvious to one having ordinary skill in the art at the time the invention was made provide Hossainy's modified implantable device such that it has adequate rigidity from about three to about six

months following implantation as taught by Igaki. Doing so would provide adequate support to a lumen until reconstriction of the lumen is no longer an issue.

With further respect to claims 7, 10, and 50, such materials are well known in the art and thus it would have been obvious to one having ordinary skill in the art at the time the invention was made to form the inner core and covering from the materials recited, since it has been held to be within the general skill of a worker in the art to select a known material on the basis of its suitability for the intended use as a matter of obvious design choice. *In re Leshin*, 125 USPQ 416.

With further respect to claims 11-13, Hossainy discloses the inner core material may comprise a cylindrical stent with perforated passages, a cylindrical structure formed of helical wound or serpentine wire structures, or a rolled tubular structure that is woven, wrapped, drilled, etched, or cut to form passages. Hossainy fails to disclose whether the filaments utilized are monofilaments or multifilaments. However, it would have been obvious to one having ordinary skill in the art at the time the invention was made to construct the inner core comprising either monofilaments or multifilaments, since such configurations are well known in the art and the applicant has failed to disclose that such configurations provide an advantage, are used for a particular purpose, or solve a stated problem. It appears the invention would perform equally well with any configuration, including configurations disclosed by Hossainy.

Claim 15 is being treated as a product by process limitation, in that "the tubular structure is micromachined or laser-cut" refers to the process of forming the tubular structure and not to the final product created. As set forth in MPEP 2113, "Even though

product-by-process claims are limited by and defined by the process, determination of patentability is based on the product itself. The patentability of a product does not depend on its method of production. If the product in the product-by-process claim is the same as or obvious from a product in the prior art, the claim is unpatentable even though the prior product was made by a different process." In re Thorpe, 777 F.2d 695,698,227 USPQ 964,966 (Fed. Cir. 1985). Examiner has evaluated the product claim without giving much weight to the method of its manufacture. Therefore, in this case, a stent as described above wherein the tubular body is formed by micromachining or laser-cutting is directed to the method of making the stent and not to the final product made. It appears that Hossainy's modified product would be the same as that claimed, especially since both applicant's product and the prior art product have the same final structure of a biodegradable inner tubular structure and a biodegradable covering material.

With further respect to claims 19-21, Hossainy's modified stent is capable of being used as claimed if one so desires.

Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Melanie Tyson whose telephone number is (571) 272-9062. The examiner can normally be reached on Monday through Thursday 8-7 (IFP).

If attempts to reach the examiner by telephone are unsuccessful, please contact the examiner's supervisor, Corrine McDermott, at (571) 272-4754. The fax phone

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number for the organization where this application or proceeding is assigned is 571-273-8300.

If there are any inquiries that are not being addressed by first contacting the Examiner or the Supervisor, you may send an email inquiry to TC3700_Workgroup_D_Inquiries@uspto.gov.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Melanie Tyson/
Primary Examiner, Art Unit 3773
July 2, 2011